Quality Improvement Organization Manual Chapter 1 - Background and Responsibilities

Table Of Contents

(Rev. 16, 06-30-06)

1000 - Authority

1005 - Purpose of QIO Review

1010 - QIO Responsibilities

1015 - Centers for Medicare & Medicaid Services (CMS) Role

1020 - Health Care Quality Improvement Program (HCQIP)

1025 - Hospital Payment Monitoring Program (HPMP)

1030 - QIO Support Center (QIOSC)

1000 – Authority

(Rev. 16, Issued: 06-30-06, Effective: 06-30-06, Implementation: 06-30-06)

Part B of title XI of the Social Security Act (the Act), as amended by the Peer Review Improvement Act of 1982, established the Utilization and Quality Control Peer Review Organization Program -- the Quality Improvement Organization (QIO) Program. The QIO program activities are further governed by title XVIII of the Act, as amended, and by regulations contained in:

- 42 CFR 405, 411 -- Limitation on liability;
- 42 CFR 412 -- Outlier review, Diagnosis Related Group (DRG) validation, and hospital notices of non-coverage;
- 42 CFR 475 -- Definition of eligible organizations and area designation;
- 42 CFR 476 -- Assumption and conduct of review;
- 42 CFR 478 -- QIO reconsiderations and appeals;
- 42 CFR 480 -- Disclosure of information;
- 42 CFR 482 -- Hospital Conditions of Participation; and
- 42 CFR 1004 -- Sanctions.

1005 - Purpose of QIO Review

(Rev. 16, Issued: 06-30-06, Effective: 06-30-06, Implementation: 06-30-06)

As a result of legislative mandates and the Centers for Medicare & Medicaid Services' (CMS) experience in administrating the program, CMS has identified the following requirements of the QIO program:

- *Improve quality of care for beneficiaries*;
- Protect the integrity of the Medicare Trust Fund by ensuring that Medicare pays only for services and goods that are reasonable and medically necessary and that are provided in the most appropriate setting; and
- Protect beneficiaries by expeditiously addressing individual complaints, notices, and appeals, such as beneficiary complaints; provider-issued notices of noncoverage (Hospital-Issued Notice of Non-coverage [HINN], Notice of Discharge and Medicare Appeal Rights [NODMAR], and Medicare Advantage appeal); Emergency Medical Treatment and Labor Act (EMTALA) violations; and other related statutory QIO responsibilities. Also, protect beneficiaries by

implementing a fast-track appeals process that limits the beneficiaries' financial liability when a Notice of Medicare Non-coverage has been issued from a provider (i.e., Skilled Nursing Facility (SNF), Home Health Agency (HHA), Hospice, Comprehensive Outpatient Rehabilitation Facility (CORF)).

Moreover, you review items or services provided to Medicare beneficiaries as instructed in the QIO Manual, Chapters 4, 5, 7, and 9.

1010 - QIO Responsibilities

(Rev. 16, Issued: 06-30-06, Effective: 06-30-06, Implementation: 06-30-06)

- A. Responsibilities Prior to Review -- You are responsible for:
 - Obtaining written Memorandums of Agreement (MOAs) (see Chapter 3);
 - Specifying in your review plan and instructions to practitioners and providers the
 type of evidence you require to document that the care ordered or furnished was
 medically necessary, reasonable, and appropriate, and that the quality of services
 met professionally recognized standards of health care; and
 - Applying professionally developed criteria for providing care, diagnosis, and treatment based upon typical patterns of practice within your geographic area to evaluate the medical necessity, quality, or appropriateness of services ordered or furnished.
- B. Ongoing Review Activities -- As a part of your ongoing review activities, you must:
 - Notify the appropriate agency of the State or Federal government when you become aware of situations which appear to be improper, but which do not fall within your review responsibilities (e.g., poor quality care in a renal dialysis center);
 - Use your authority or influence to enlist the support of other professional or government agencies to ensure that all providers and practitioners for which you have review responsibilities comply with their obligations (see §1156 of the Act.); and
 - Conduct beneficiary outreach and education activities for the express purpose of informing beneficiaries about:
 - o The QIO program and how to contact the QIO;
 - o Beneficiary rights as outlined at §1154(a)(4)(B) of the Act; and
 - o How to exercise those rights, including what to expect when they do contact the QIO (e.g., length of time to obtain a response, form the

response will take). This information must include the processes regarding beneficiary complaints and notices of non-coverage (e.g., *HINN* and *NODMAR*).

- C. Responsibilities as a Result of Your Review -- To act upon information you obtain as a result of your review activities, you must:
 - At least annually, publish and distribute to providers and practitioners, whose services you review, a report of your activities and findings as required in §1154(a)(6)(B) of the Act. It must include:
 - A description of the types of cases where you have found inappropriate or unnecessary care, services that were rendered in an inappropriate setting, and/or services that did not meet professionally recognized standards of care; and
 - A description of your Health Care Quality Improvement Program (HCQIP) activities.
 - Identify and seek correction of situations that, if continued, would result in violations under §1156 of the Act. This includes referring certain cases to state licensing boards (see §1154(a)(9)(B) of the Act);
 - Submit reports to the Office of the Inspector General (OIG) on providers and practitioners found to have substantially violated an obligation in a substantial number of cases, or to have grossly and flagrantly violated an obligation in one or more instances; and
 - Coordinate your activities, including information exchanges, in order to promote the efficient and economical operation of programs among appropriate public and private agencies. This fulfills §1154(a)(10) of the Act, but is subject to confidentiality rules in §1160 of the Act and regulations at 42 CFR Part 480. It includes, at a minimum:
 - o Meeting with the State Agencies; and
 - O Communicating with accrediting bodies, quality organizations, and any other agencies as necessary to carry out QIO activities.
- D. Additional Responsibilities -- Perform all other activities specified in the Statement of Work (SOW) of your CMS contract, including any modifications, CMS regulations and instructions, and relevant statutory provisions. The SOW includes your requirements for improving quality of care for beneficiaries and protecting beneficiaries and the Medicare program.

1015 - Centers for Medicare & Medicaid Services (CMS) Role (Rev. 10, 09-05-03)

The CMS was established in March 1977, to combine health care financing and quality assurance programs into a single agency. CMS is responsible for the Medicare program, Federal participation in the Medicaid program, the Quality Improvement Organization (QIO) program, and a variety of other health care quality assurance programs.

A. The Primary Mission

The CMS' primary mission is to administer its programs in a manner that:

- > Promotes the timely delivery of appropriate, quality health care to beneficiaries;
- Ensures that beneficiaries are aware of the services for which they are eligible;
- Ensures that those services are accessible and of high quality; and
- > Promotes efficiency and quality within the total health care delivery system.
- B. Central Office Policy-making Responsibility -- Overall policy-making responsibility for administration of the QIO program is centralized in CMS' Office of Clinical Standards and Quality (OCSQ). OCSQ is responsible for:
 - Monitoring and overall administrative control of the QIO program, including coordinating with CMS' Office of Internal Customer Support on contracts and financial aspects;
 - Establishing operational policy for the QIO program; and
 - ➤ Developing operational instructions and official interpretations of policy for QIOs and CMS Regional Offices (ROs).

C. Regional Office Assistance to QIOs

The ROs are responsible for assuring that QIOs meet applicable Federal requirements under the provisions of their contracts. The ROs:

- ➤ Provide liaison, direction, and technical assistance to QIOs in the day-to-day management of their operations;
- > Interpret CMS guidelines, policies, and procedures applicable to QIO activities;
- ➤ Analyze QIO budgets and spending patterns to assure that funds are economically and appropriately utilized;

- ➤ Recommend the allocation of funds for conducting additional activities;
- > Conduct assessments of QIO operations;
- ➤ Review QIO actions; and
- Provide feedback to each QIO.

1020 - Health Care Quality Improvement Program (HCQIP)

(Rev. 16, Issued: 06-30-06, Effective: 06-30-06, Implementation: 06-30-06)

The Health Care Quality Improvement Program (HCQIP) had its conceptual foundations in the health care variations research of the 1980s that examined variability in care and outcomes among providers and geographic areas. Under HCQIP, you conduct Quality Improvement Projects (QIPs) using statistical information to examine medical processes and outcomes of health care, and provide feedback to providers and practitioners so that this information can be used to benchmark progress toward improved practices and outcomes. CMS designed HCQIP to improve health outcomes of all Medicare beneficiaries regardless of personal characteristics (e.g., socio-economic status, health status, ethnic group), physical location (urban or rural), or setting (e.g., physicians' offices, Medicare Advantage Organizations, hospitals, nursing homes). Your SOW sets specific quality measures for national health improvement priorities that reflect the current state of QIO program experience, measurement systems, and data sources. These quality measures do not address the entire spectrum of health care, nor do they reflect fully the unique circumstances of each state (see Chapter 16).

1025 - Hospital Payment Monitoring Program (HPMP)

(Rev. 16, Issued: 06-30-06, Effective: 06-30-06, Implementation: 06-30-06)

OIG Audit Opinion of CMS' 1997 Financial Statement found that approximately \$4 billion in improper payments were made for inpatient services under the Prospective Payment System (PPS). In order to reduce this payment error rate, you must initiate a program of projects. CMS defines the payment error rate as the number of dollars found to be paid in error out of the total of all dollars paid for inpatient PPS services. CMS will implement a surveillance system to provide state-specific estimates of the payment error rate. These estimates will be used as performance indicators on which to evaluate your performance.

CMS has developed the Comprehensive Plan for Program Integrity (Plan) to serve as the road map for reducing payment errors in the Medicare program. The Hospital Payment Monitoring Program (HPMP, formerly the Payment Error Prevention Program or PEPP) is one of ten initiatives included in this Plan.

This ongoing surveillance allows CMS to continue to produce accurate and reliable estimates of payment errors at the State and national level for inpatient acute care

hospital PPS services. In addition, highly targeted, narrowly focused projects will be undertaken by some QIOs in an effort to reduce known sources of payment error.

1030 - QIO Support Center (QIOSC)

(Rev. 16, Issued: 06-30-06, Effective: 06-30-06, Implementation: 06-30-06)

For certain topics, settings, populations, and project processes, CMS is contracting with designated QIOs to provide support for CMS and the QIO community for the particular areas of interest. These QIO Support Centers (QIOSCs) will work in conjunction with corresponding Topic Area Teams (TATs) established by CMS. The QIOSC will serve as a clearinghouse of topic-specific information and provide technical assistance to QIOs as necessary.